

Research Article

A new gorgonian *Pseudopterogorgia nanjiensis* sp. nov. (Cnidaria, Octocorallia, Gorgoniidae) from the Nanji Islands, China

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Abstract

Members of the genus *Pseudopterogorgia* Kükenthal, 1919 are widely distributed in shallow water of the Indo-West Pacific. During an investigation of benthic biodiversity in the subtidal zone surrounding the Nanji Islands in the East China Sea, two specimens of *Pseudopterogorgia* were collected and described as a new species based on an integrated morphological-molecular approach. *Pseudopterogorgia nanjiensis* **sp. nov.** is most similar to *P. fredericki* Williams & Vennam, 2001 in the irregular branching form and indistinct scaphoids, but differs by the coenenchymal sclerite content of distinct capstans and a few warty spindles and radiates (vs. mostly warty spindles and a few capstans), and a purplish colony (vs. white, pink to deep rose). Molecular phylogenetic analyses, based on the mtMutS-COI gene sequences, delineated a monophyletic clade encompassing all assessed *Pseudopterogorgia* species. Within this clade, *P. nanjiensis* **sp. nov.** showed a close phylogenetic affinity with both *P. fredericki* and *P. australiensis* Ridley, 1884.

Key words: Anthozoa, COI, Malacalcyonacea, morphology, mtMutS, new species, phylogeny, *Pseudopterogorgia nanjiensis*, taxonomy, 28S rDNA



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Introduction

Octocorals fulfill a pivotal ecological function, with their three-dimensional configurations enhancing the structural intricacy and variety of reefs (Poliseno 2016; Lau et al. 2020). Gorgonian corals in octocoral animal forests offer nourishment and habitat for a variety of interrelated organisms (Sánchez 2016). Among these, the genus *Pseudopterogorgia* Kükenthal, 1919 represents a rare group of shallow-water octocorals found in the Indo-West Pacific (Williams and Chen 2012). The members of this genus are characterized by pinnate or irregular branching structures, absence of anastomoses, and C-shaped or scaphoid-like sclerites present in the coenenchyme (Kükenthal 1919; Bayer 1961; Grasshoff and Alderslade 1997). To date, this genus has seven valid species, all

of which are distributed in the Indo-West Pacific (Williams and Chen 2012; WoRMS Editorial Board 2024).

During the investigation of benthic biodiversity in the subtidal zone of the Nanji Islands on the Chinese coast of the East China Sea, two specimens of *Pseudopterogorgia* were collected. They are described as a new species *Pseudopterogorgia* nanjiensis sp. nov. based on morphological and phylogenetic analyses. The phylogenetic placement of *P. nanjiensis* sp. nov. within the genus is further explored.

Material and methods

Specimen collection and morphological examination

Two specimens were obtained by scuba diving from the subtidal zone of the Nanji Islands (27°28.53'N, 121°08.13'E) on the Chinese coast of the East China Sea during investigation of the benthic biodiversity in 2022 and 2023 (Fig. 1). The specimens were photographed in situ using a Nikon D850 before being sampled and on board using a Canon EOS 5D Mark IV, and then stored in 75% ethanol after collection.

A stereo dissecting microscope (Zeiss SteREO Discovery. V20) was used to examine the general morphology and anatomy of the specimens. Sclerites of the polyps and branches were isolated by digesting the tissues in sodium hypochlorite and subsequently rinsed thoroughly with deionized water. To investigate the ultrastructure of polyps and sclerites, they were air-dried, mounted on carbon double adhesive tape, and coated with Pd/Au for scanning electron microscope (SEM) observation. SEM scans were obtained by using a TM-3030Plus SEM at 15 kV, and the optimum magnification was selected for each type of sclerite. The terminology used in this study follows Bayer et al. (1983).

The type specimens of the new species have been deposited in the Marine Biological Museum of Chinese Academy of Sciences (MBMCAS) at Institute of Oceanology, Chinese Academy of Sciences, Qingdao, China.

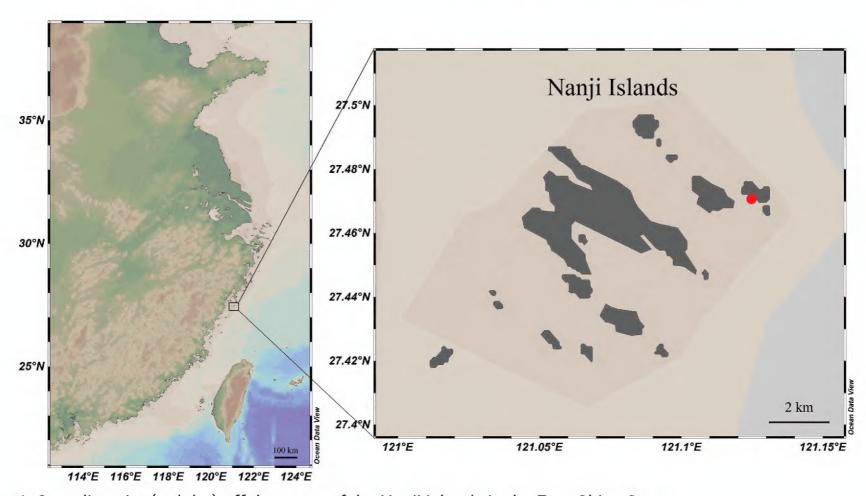


Figure 1. Sampling site (red dot) off the coast of the Nanji Islands in the East China Sea.

DNA extraction and sequencing

Genomic DNA was extracted from the polyps of each specimen using the DNeasy Blood and Tissue Kit (Qiagen, Hilden, Germany) following the instructions. Specific regions of the mitochondrial *mutS* homolog (mtMutS) and cytochrome *c* oxidase subunit I (COI), as well as an approximately 800-nt fragment of the 28S nuclear ribosomal gene (28S rDNA), were selected for phylogenetic analysis. For the amplification of mtMutS and COI, the primer pairs AnthoCorMSH (5'-AGGAGAATTATTCTAAGTATGG-3'; Herrera et al. 2010) and Mut-3458R (5'-TSGAGCAAAAGCCACTCC-3'; Sánchez et al. 2003); COI8414-F (5'-CCAGGTAGTATGTTAGGRGA-3') and HCO2198 (5'-TAAACTTCAGGGTGACCAAAAAATCA-3') were utilized, respectively. Sequencing of 28S rDNA was performed using the primers 28S-Far (5'-CACGAGACCGATAGCGAACAAGTA-3') and 28S-Rar (5'-TCATTTCGACCCTAAGACCTC-3') (McFadden and van Ofwegen 2012). Amplifications and sequencing of the markers followed Xu et al. (2021).

Genetic distance and phylogenetic analyses

The sequences of Gorgoniidae species and two species of Eunicellidae as an out-group were downloaded from GenBank (Table 1). The sequences were aligned using MAFFT v.7 (Katoh and Standley 2013) with the L-INS-I algorithm. Genetic distances of mtMutS, COI and 28S between species/populations were calculated with MEGA 6.0 using Kimura 2-parameter model (Tamura et al. 2013).

Table 1. The sequences used in this study. Newly sequenced species are in bold.

| Smaning | Voucher Number | GenBank Accession Numbers | | | |
|---------------------------------------|----------------------------------|---|----------|----------|--|
| Species | voucher Number | mtMutS | COI | 28S rDNA | |
| Pseudopterogorgia nanjiensis sp. nov. | MBM287892 | PP558212 | PP556906 | PP572455 | |
| Pseudopterogorgia nanjiensis sp. nov. | MBM287893 | PP558213 | PP556907 | PP572456 | |
| Pseudopterogorgia australiensis | - | AY268442 | _ | _ | |
| Pseudopterogorgia fredericki | CAS:118507 | JX152766 | _ | _ | |
| Pseudopterogorgia rubrotincta | CAS:155043 | JX152768 | _ | _ | |
| Leptogorgia alba | -/HMG71/MECN Ant0018 | AY268452 | HG917083 | KX721241 | |
| Leptogorgia cuspidata | -/HMG97/MCZ:7002 | -/HMG97/MCZ:7002 AY268450 HG917088 | | KX767433 | |
| Leptogorgia mariarosae | MECN Ant0012 | KX721193 | KX721174 | KX721231 | |
| Leptogorgia obscura | Ant 0066/Ant 0066/BEIM 0080 | Ant 0066/Ant 0066/BEIM 0080 KX767321 KX767383 | | KX721248 | |
| Leptogorgia cf. palma | USNM:1516865/- | ON109715 | MW401657 | _ | |
| Eugorgia daniana | MECN Ant0033 | 33 KX721208 KX72118 | | KX721246 | |
| Pacifigorgia irene | Ant 0044/Ant 0044/MNCN 2.04/1174 | KX767346 | KX767406 | KX767449 | |
| Gorgonia flabellum | -/-/RMNH Coel.40827 | AY126427 | GQ342418 | JX203708 | |
| Phyllogorgia dilatata | - | AY126428 | <u>-</u> | _ | |
| Antillogorgia bipinnata | -/ABBAC034/RMNH Coel.40828 | GQ342499 | MK153463 | JX203712 | |
| Adelogorgia osculabunda | MZUCR:2496_OCT0088 | MF579541 | _ | _ | |
| Psammogorgia cf. arbuscula | HMG38/HMG15/HMG38 | G38 HG917043 HG917055 | | LT221092 | |
| Chromoplexaura marki | - | KX904972 | KX904954 | _ | |
| Callistephanus sp. | USNM:1606569 | ON109735 | - | _ | |
| Eunicella tricoronata | RMNH Coel.40814 | JX203795 | JX203853 | JX203707 | |
| Complexum monodi | CSM-SEN3/CSM-SEN4/CSM-SEN4 | KP036906 | KP036909 | KP036912 | |

Phylogenetic analyses were conducted on the 28S rDNA and mtMutS-COI datasets. The phylogenetic frameworks were established using PhyloSuite v.1.2.3 (Zhang et al. 2020; accessible at http://phylosuite.jushengwu.com), a comprehensive platform for phylogenetic analysis. ModelFinder v.2.2.0 (Kalyaanamoorthy et al. 2017) was employed to identify the optimal evolutionary models based on the Bayesian Information Criterion (BIC). HKY+F+I and TIM3e+I were selected as the best-fit substitution models for the mt-MutS-COI and 28S rDNA alignments, respectively. Maximum likelihood (ML) trees were inferred using IQ-TREE v.2.2.0 (Nguyen et al. 2015) with 1000 standard bootstraps. Following Hillis and Bull (1993), ML bootstrap values were categorized into low (<70%), moderate (70-90%), and high ($\ge90\%$) confidence levels. Bayesian inference (BI) phylogenies were reconstructed using MrBayes v.3.2.7a (Ronquist et al. 2012) with dual parallel runs, each consisting of 10,000,000 generations and a burn-in phase that discarded the initial 25% of sampled data. Following Alfaro et al. (2003), the Bayesian posterior probabilities < 0.95 and ≥ 0.95 were considered as low and high, respectively.

Results

Phylum Cnidaria Hatschek, 1888
Subphylum Anthozoa Ehrenberg, 1834
Class Octocorallia Haeckel, 1866
Order Malacalcyonacea McFadden, van Ofwegen & Quattrini, 2022
Family Gorgoniidae Lamouroux, 1812
Genus *Pseudopterogorgia* Kükenthal, 1919

Pseudopterogorgia nanjiensis Sun, Xu & Zhan, sp. nov. https://zoobank.org/37902653-255A-46C8-9867-7C8DDC045BA3 Figs 2, 3, Table 2

Material examined. *Holotype*: MBM287893, CHINA-Zhejiang Prov. • Nanji Islands; 27°28.53'N, 121°08.13'E; 11 May 2023. *Paratype*: MBM287892, CHINA-Zhejiang Prov. • Nanji Islands; 27°28.53'N, 121°08.13'E; 14 m; 11 May 2022.

Diagnosis. Colony non-pinnate and purplish. Branches divided dichotomously and irregularly, nearly in one plane. Polyp retracted and forming a small and oval-shaped protrusion. Sclerites in polyps small flat rodlets, colorless and rare to absent in numbers. Sclerites in coenenchyme red, mostly capstans, a few radiates and warty spindles, and rare scaphoids. Capstans have two whorls of tubercles and blunt ends with irregular arrangements of complex tubercles. Radiates and immature sclerites with two whorls of projections, spindles with 4–8 whorls of tubercles, and scaphoids with similar ornamentation of tubercles on both convex and concave sides.

Description. Holotype upright and nearly planar but not pinnate, about 70 mm in length (Fig. 2B). Holdfast nearly round, about 8 mm long and 7 mm wide at maximum. Trunk cylindrical, about 3 mm in length before first branch and 2 mm in diameter at base, divided into two primary stems, one with three secondary branches and the other with five secondary branches. Branches divided dichotomously and irregularly, nearly in one plane (Fig. 2D). Distances between

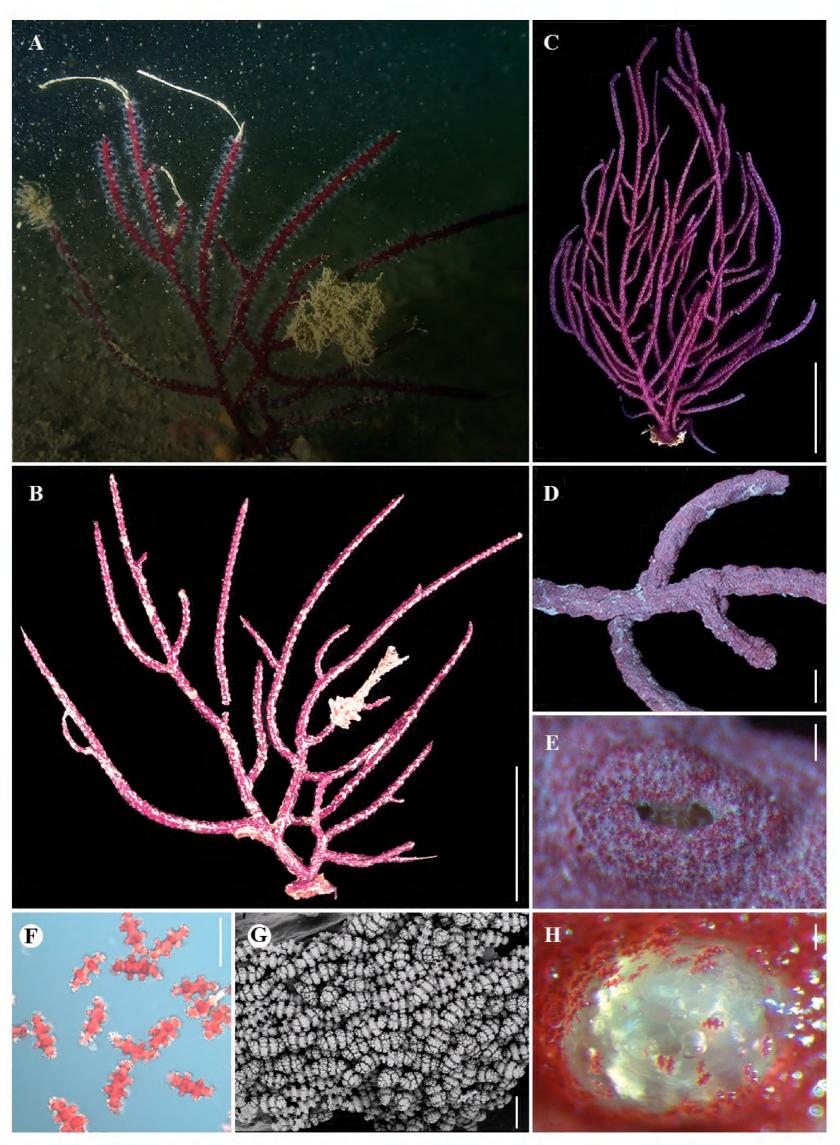


Figure 2. The external morphology and polyps of *Pseudopterogorgia nanjiensis* sp. nov. A holotype in situ **B** holotype after collection **C** paratype after fixation **D** part of branch under a light microscope **E** single polyp aperture under a light microscope **F** sclerites under a light microscope **G** coenenchyme under SEM **H** single dissolved polyp aperture under a light microscope. Scale bars: 2 cm (**B**); 3 cm (**C**); 2 mm (**D**); 0.2 mm (**E**); 0.1 mm (**F**, **G**, **H**).

adjacent branches up to 20 mm. Branches usually cylindrical in the proximal part of the colony, and became flattened in the distal part of the colony. Terminal branchlets slender and a little curved, typically measuring 20–60 mm in length and 1–2 mm in width.

Table 2. Comparison of morphological characters of *Pseudopterogorgia* species (based on Williams and Vennam 2001). "-" means unavailable data.

| Characters/ Species | P. nanjiensis sp. nov. | P. fredericki | P. formosa | P. australiensis | P. luzonica | P. oppositipinna | P. rubrotincta | P. torresia |
|-------------------------------|---|---|---|--|---|-------------------------------------|--|---|
| Branching | irregular | irregular | irregular | pinnate/ plumose | lateral | pinnate to irregular | pinnate | pinnate/ sparse |
| Sclerite forms | capstans, radiates, warty spindles, scaphoids, flat rodlets | elongated spindles, some shorter and more robust spindles, scaphoids | small or double or girdled spindles, a few double heads, scaphoids | straight or slightly curved spindles, eight radiates, scaphoids | girdled spindles, scaphoids | elongated spindles, scaphoids | rough spindles, scaphoids | warty spindles, quadriradiate, scaphoids |
| Scaphoids | indistinct | indistinct | some distinct | some distinct | mostly distinct | mostly distinct | some distinct | some distinct |
| Colony color | purplish | white, pink to deep rose | dark bright pink to carmine | yellow, orange or deep crimson | bright red | brown to amber-yellow. | orange yellow with red line | pale yellow |
| Sclerites color | red or colorless | reddish or colorless | red, carmine | yellow or red | red | purple-red or crimson red | red | _ |
| Maximum sclerite length | 0.18 mm | 0.23 mm | 0.17 mm | 0.17 mm | 0.16 mm | 0.21 mm | 0.15 mm | 0.16 mm |
| Type locality | Nanji Islands on East China Sea | western India | Lombok | Torres Strait | Luzon | Mergui Archipelago | Sri Lanka | Torres Strait |
| Distribution | Western Pacific | Indian Ocean | Western Pacific | Western Pacific | Western Pacific | Indian Ocean | Indian Ocean | Western Pacific |
| Depth (m) | 14 | 6-8 | 22 | 12-37 | _ | _ | - | _ |
| References | present study | Williams and Vennam 2001; Williams and Chen 2012; Ramvilas 2023 | Nutting 1910; Grasshoff and Alderslade 1997 | Ridley 1884; Thomson and Henderson 1905; Kükenthal 1919, 1924; Williams and Vennam 2001 | Kükenthal 1919; Grasshoff and Alderslade 1997 | Ridley 1888; Kükenthal 1924 | Thomson and Henderson 1905; Williams and Vennam 2001; Williams and Chen 2012; Ramvilas 2023 | Wright and Studer 1889; Williams and Vennam 2001 |

Polyp retracted, forming small and oval-shaped protrusions with a recessed orifice (Fig. 2E), measuring on average 1.00 mm in length and 0.65 mm in width. Each polyp's orifice appears as a tiny and pale yellowish slit aligned lengthwise. Polyps usually closely and irregularly spaced on the branches and sparse at the basal part of the colony, some of them arranged into two opposing longitudinal rows on the terminal branchlets. Distance between polyps no more than 4 mm.

Coenenchyme covered with dense sclerites, while the polyps sclerites rare to absent (Fig. 2G, H). Sclerites in coenenchyme red with various shapes, arranged in two inconspicuous layers without a clear boundary, including abundant capstans, rare indistinct scaphoids, and occasional tuberculate spheroids and crosses in outer layers, a few radiates and warty spindles in inner layers. These sclerites range in length from 0.05 mm to 0.18 mm, but mostly less than 0.12 mm long (Figs 2F, 3A–E). Among them, capstans have two whorls of tubercles, up to 0.13 mm long, and blunt ends with irregular arrangements of complex tubercles (Fig. 3A). The radiates and immature sclerites with two whorls of projections and often blunt ends, some of them covered with shallow longitudinal grooves on surface, measuring 0.05–0.09 mm (Fig. 3B). The warty spindles with 4 to 8 transversely-aligned whorls of tubercles, up to 0.18 mm in length (Fig. 3C). The scaphoids, not distinctly developed and resembling curved spindles, with 4–8 noticeable whorls of tubercles, up to 0.14 mm long

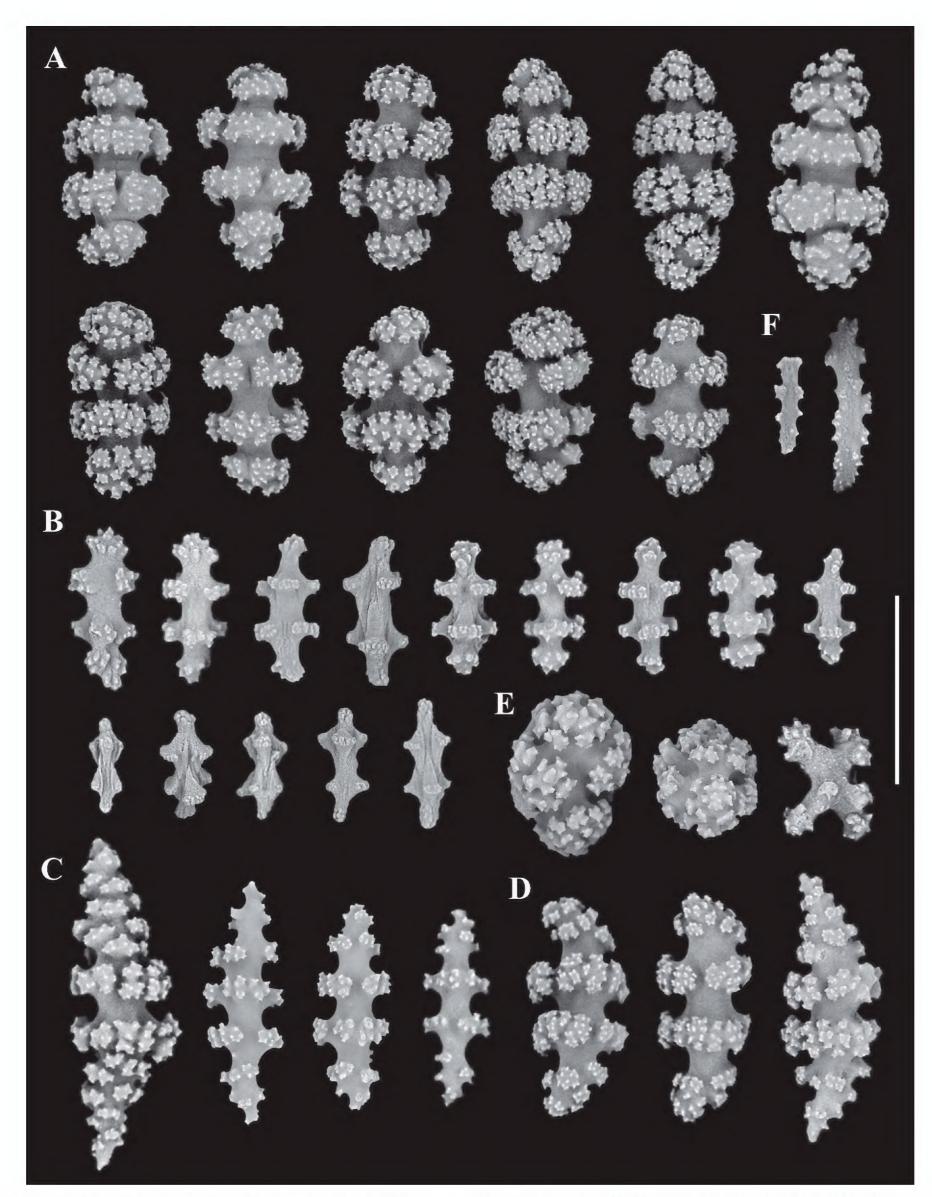


Figure 3. The sclerites of *Pseudopterogorgia nanjiensis* sp. nov. **A** capstans **B** radiates and immature sclerites **C** warty spindles **D** scaphoids **E** tuberculate spheroids and cross **F** flat rodlets. Scale bars: 0.1 mm (all in the same scale).

(Fig. 3D). The convex and concave sides have a similar degree of fine ornamentation on the whorls of tubercles, and there seems to be minimal variation in the tubercles between the two sides with those on the convex sides potentially being slightly shorter. Tuberculate spheroids and crosses occasionally present

in coenenchyme, are up to 0.08 mm and 0.06 mm long, respectively (Fig. 3E). Small rodlets in polyps flat and colorless, rare in numbers and only found in a few polyps, covered with sparse warts on the edges, up to 0.09 mm (Fig. 3F).

Holotype purplish in situ and after collection, and became deep purplish red after fixation.

Variation of paratypes. The two specimens exhibited a high degree of morphological concordance, with discrepancies primarily noted in colony size and branching density. Paratype irregularly branched, about 140 mm in length and 90 mm in width (Fig. 2C). The main trunk before first branch about 5 mm in length and 2 mm in diameter at base. The terminal branchlets 30–100 mm long and 1–2 mm wide. The oval-shaped protrusions formed by retracted polyps about 1.00 mm long and 0.75 mm wide. The warty spindles in coenenchyme up to 0.16 mm long.

Type locality. The subtidal zone of Nanji Islands with water depth of 14 m. **Etymology.** Named after the type locality Nanji Islands.

Distribution and habitat. Known only from the subtidal zone of the Nanji Islands on the Chinese coast of the East China Sea with a water depth of 14 m. Colony attached to a rocky substrate. The water temperature was 18 °C and the pH was 8.13.

Remarks. Given the non-uniform nature of these variations, they are treated as manifestations of intraspecific variability. *Pseudopterogorgia nanjiensis* sp. nov. is most similar to *P. fredericki* Williams & Vennam, 2001 in the irregular branching form and indistinct scaphoids, but differs by the sclerite forms and sizes (almost capstans, a few warty spindles and radiates, and rare indistinct scaphoids and small rodlets, mostly less than 0.12 mm vs. almost warty spindles and a few scaphoids, mostly more than 0.12 mm) and the purplish colony (vs. white, pink to deep rose) (Williams and Vennam 2001; Figs 2, 3, Table 2). *Pseudopterogorgia nanjiensis* sp. nov. is also analogous to *P. formosa* Nutting, 1910 in the irregular branching form, but distinct from *P. formosa* by the sclerite forms (capstans, spindles, radiates, and scaphoids vs. spindles, double heads, and scaphoids) (Nutting 1910; Table 2). In addition, the new species can be readily distinguished from the remaining congeners by its irregular branching form (vs. pinnate or lateral; Table 2).

Genetic distance and phylogenetic analyses

The new sequences were deposited in GenBank (Table 1). The alignments comprised 720, 575 and 717 nucleotide positions for the mtMutS, COI, and 28S rDNA regions, respectively. Based on the aligned region of mtMutS, the interspecific distance of *P. australiensis* Ridley, 1884, *P. fredericki*, *P. nanjiensis* sp. nov. and *P. rubrotincta* Thomson & Henderson, 1905 ranged from zero to 0.44%, while the intraspecific distance was zero, which was calculated from only two specimens of *P. nanjiensis* sp. nov. (Suppl. material 1: table S1). The mtMutS genetic distances between *P. nanjiensis* sp. nov. and congeners are in the range of 0–0.28%. For the 28S rDNA and COI regions, only sequences of *P. nanjiensis* sp. nov. are currently available within the genus *Pseudopterogorgia*, and no genetic variation was detected between the two specimens analyzed (Suppl. material 1: tables S2, S3). The genetic distances between *Pseudopterogorgia* and other genera of Gorgoniidae are greater than 5.48% for mtMutS, 5.74% for 28S, and 2.04% for COI (Suppl. material 1: tables S1–S3).

The Bayesian inference (BI) tree is nearly identical to the maximum likelihood (ML) tree in topology for both the mtMutS-COI and 28S rDNA regions, and thus only the BI tree annotated with support values from both inference methods is presented (Figs 4, 5). In the mtMutS-COI trees, all the *Pseudopterogorgia* species formed a monophyletic clade with full node support. The new species *P. nanjiensis* sp. nov. clustered with the clade comprising *P. fredericki* and *P. australiensis* (Fig. 4). In the 28S rDNA trees, *P. nanjiensis* sp. nov. emerged as a distinct clade, with a basal branching position within the family Gorgoniidae (Fig. 5).

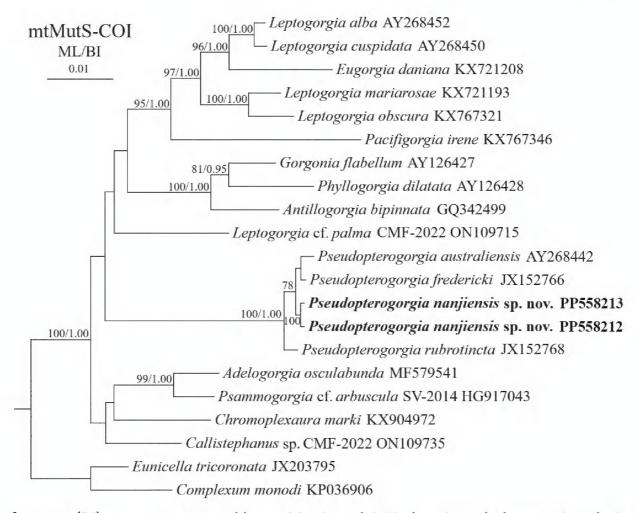


Figure 4. Bayesian inference (BI) tree constructed by mtMutS and COI showing phylogenetic relationships among *Pseudopterogorgia* and related genera and species. The maximum likelihood (ML) tree is identical to the BI tree in topology. Node support is as follows: ML bootstrap/BI posterior probability. The ML bootstrap < 70% or BI posterior probability < 0.95 is not shown. Newly sequenced species are in bold. The GenBank accession numbers of the mtMuts sequences are listed next to the species names.

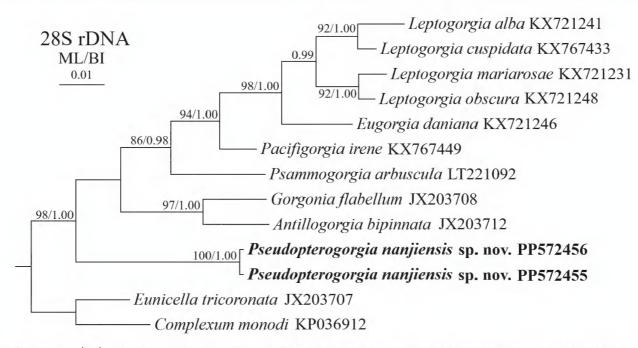


Figure 5. Bayesian inference (BI) tree constructed by 28S rDNA showing phylogenetic relationships among *Pseudopterogorgia* and related genera and species. The maximum likelihood (ML) tree is identical to the BI tree in topology. Node support is as follows: ML bootstrap/BI posterior probability. The ML bootstrap < 70% or BI posterior probability < 0.95 is not shown. Newly sequenced species are in bold. The GenBank accession numbers of the 28S rDNA sequences are listed next to the species names.

Discussion

The genus *Pseudopterogorgia* is distinguished by its pinnate or irregular branching structures, absence of anastomoses, and C-shaped or scaphoid-like sclerites present in the coenenchyme (Kükenthal 1919; Bayer 1961; Grasshoff and Alderslade 1997). Based on the branching structure and sclerite form, the present species is undoubtedly assigned to this genus. In the mtMutS-COI trees, *Pseudopterogorgia nanjiensis* sp. nov. showed a phylogenetic affinity with *P. fredericki* and *P. australiensis*, congruent with morphological assessments (Fig. 4). All three species are characterized by the absence of anastomoses and the presence of scaphoid-like sclerites within the coenenchyme. However, *P. nanjiensis* sp. nov. can be differentiated from *P. fredericki* based on sclerite morphology (see above remarks), and it is easily separated from *P. australiensis* by the branching form (irregular vs. pinnate/plumose; Table 2; Ridley 1884; Williams and Vennam 2001). With the addition of the new species *P. nanjiensis* sp. nov. discovered on the Nanji Islands, there are currently a total of eight valid species in *Pseudopterogorgia* (Williams and Vennam 2001; WoRMS Editorial Board 2024).

In the present study, the precise phylogenetic placement of the genus *Pseudopterogorgia* remains vague, primarily attributed to the low support value assigned to the corresponding node in the mtMutS-COI trees (Fig. 4). Although the 28S rDNA trees exhibit high support values for the majority of internal branches, the systematic position of *Pseudopterogorgia* remains unresolved (Fig. 5). This uncertainty arises from the inclusion of sequences representing only seven out of a total of thirteen genera currently recognized within the family Gorgoniidae, and is further compounded by the acknowledged reality that many of these genera exhibit polyphyletic or paraphyletic relationships (Vargas et al. 2014; Poliseno et al. 2017). Therefore, to achieve a clearer resolution, further genomic data are essential. These data should encompass a more exhaustive collection of 28S rDNA sequences and potentially include ultra-conserved elements plus exon (UCEs+exon).

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Additional information

Conflict of interest

The authors have declared that no competing interests exist.

Ethical statement

No ethical statement was reported.

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Author contributions

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Data availability

All of the data that support the findings of this study are available in the main text or Supplementary Information.

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Supplementary material 1

Genetic distance analyses

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Data type: docx

Explanation note: **table S1.** Interspecific and intraspecific uncorrected pairwise distances at mtMutS of *Pseudopterogorgia* and species of other genera within Gorgoniidae. **table S2.** Interspecific and intraspecific distances at 28S of *Pseudopterogorgia* and species of other genera within Gorgoniidae. **table S3.** Interspecific and intraspecific distances at COI of *Pseudopterogorgia* and species of other genera within Gorgoniidae.

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